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*Original Citation:*

*Availability:*

This version is available <http://hdl.handle.net/2318/144927> since 2016-10-20T16:28:53Z

*Published version:*

DOI:10.1016/j.autrev.2012.11.009

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## **Cost-effectiveness of biologic treatment for rheumatoid arthritis in clinical practice: An achievable target?**

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### **Abstract**

The burden of illness of rheumatoid arthritis (RA) falls on patients, families and society through the direct costs, indirect costs, and intangible costs. A large number of RA cost-of-illness studies have been performed in recent decades with discrepant results due to patient heterogeneity, and different health-care organization, employment rate or social support, job opportunities, and methodologies used to calculate the costs. The greatest burden of RA is the indirect and the intangible costs, but how to estimate them remains controversial. The systematic use of traditional disease modifying anti rheumatic drugs has changed the evolution of the disease. However, a considerable improvement in the management of RA has been obtained since the advent of biologic response modifiers. The use of these drugs, which have demonstrated greater efficacy than conventional therapies, have tripled the direct costs of RA, which rose from about € 4000 to roughly € 12,000, in a period of five years, from 2000 to 2005. The present paper is aimed to examine the effects of this change in therapeutic strategy.

Rheumatoid arthritis (RA) is a chronic, systemic autoimmune inflammatory arthritis that affects approximately 1% of the population [1] and [2]. The course of RA varies but most patients present symmetric polyarthritis, clinically manifesting joint pain, stiffness and swelling. Untreated, most patients have a progressive course resulting in persistent pain, progressive joint destruction and short- and long-term disability. Disease activity is predominantly responsible for the disability in the early stages of RA, while non reversible joint damage increases disability later in the course of RA. In addition, patients with RA have increased mortality compared with the general population and this is frequently attributed to an increased risk of cardiovascular disease [3], [4] and [5].

The burden of illness of RA falls on patients, families and society through the direct costs (pharmacological, surgical and rehabilitative treatment, radiology, medical appointments, hospitalization, complementary and alternative medicine, transportation reimbursement, out-of-pocket expenses for caregivers), indirect costs (loss of work productivity) and intangible costs (cost of suffering with deterioration in the quality of life of patients, their families and friends) [6], [7], [8], [9], [10], [11] and [12]. The intangible costs are difficult to quantify and are not normally considered in studies on the cost of illness. A large number of RA cost-of-illness studies have been performed in recent decades with very different results. The discrepancies in the results probably come from the heterogeneity of the patient population (number of patients, differences in severity and duration of illness), different health-care organizations, different employment rate and social support, different job opportunities and different methodologies used to calculate the costs (friction cost method, human capital approaches, payers' perspective, social perspective) [11].

From the literature it is clear that the overall mean costs of RA amount to about €15,000 per year, while the direct annual costs of RA are on average about € 4000. However, the greatest burden of RA costs is the indirect costs, although consensus is lacking as to how to estimate them [10]. Different patient characteristics have an important influence on the costs of the illness. Indeed, the direct costs increase very significantly among the four functional classes of the American College of Rheumatology [6] and [8] and the cost increases in relation to the severity of the disease [13], [14] and [15]. There is also further evidence that higher scores on the health assessment questionnaire (HAQ)—the most commonly used tool for measuring the degree of disability—corresponds to increased worsening of the disease in the future, to greater morbidity and mortality, and to a progressive increase in the direct and indirect costs [11], [16], [17] and [18]. Comorbidity—above all cardiovascular disease, infections and depression has a heavy impact on disability [19], [20] and [21] resulting in an increase in costs, especially in patients with severe, long-standing disease [6], [11], [22] and [23].

Undoubtedly, the systematic use of traditional disease modifying anti rheumatic drugs (DMARDs) such as methotrexate, leflunomide and sulfasalazine has changed the evolution of the disease. Indeed, evidence indicates that the early use of DMARDs slows both joint erosion and progression of joint damage, and preserves function [24], [25], [26], [27], [28] and [29]. Non steroidal anti-inflammatory drugs (NSAIDs) and corticosteroids can reduce inflammation, control symptoms and lessen the functional limitations caused by disease activity. Moreover, there is evidence that if corticosteroids are used in association with DMARDs they can contribute towards slowing the progression of radiological lesions [30], [31], [32] and [33].

However, a considerable improvement in the management of RA has been obtained since the advent of biologic response modifiers. Nowadays, it is realistically possible to set clinical and radiological remission of the disease as the treatment objective and maintain these conditions over time. Indeed, these drugs can swiftly control the inflammatory processes and symptoms of the disease, thus improving the quality of life and inhibiting radiological alterations and anatomical damage, thereby enabling the patient to recover and maintain joint function [28], [34], [35] and [36]. The use of these drugs, which have demonstrated greater efficacy than conventional DMARDs in controlling the disease, have led to a 4-fold to 6-fold increase in the direct costs of RA [11] and [37], since it has been estimated that their annual cost ranges between US \$16,000 and \$20,000 compared to the approximately \$3000 expense for the traditional DMARDs [38].

This has meant that in the last decade the cost of treating RA has progressively increased, above all due to the increased costs of treatment [39]. The findings of the PACTIS study in which patients took only traditional DMARDs [8] are paradigmatic compared to those of the Eco-PR study [13] where 20% of patients were treated with biologics. In a period of five years, from 2000 to 2005, the direct annual costs of RA tripled, rising from about €4000 to roughly €12,000.

This marked increase in costs has given rise to considerable anxiety in the national health agencies, and the cost-effectiveness of biologic drugs has become one of the highest priorities of health economics studies of RA. In the last decade numerous health economics surveys have been published on the use of anti TNF  $\alpha$  biologic drugs in RA. Some years ago, an overview and a review of eight pharmacoeconomics studies on anti TNF  $\alpha$  drugs [40] and [41] were presented, and very recently a systematic study was

published [39] on the cost-effectiveness of biologic drugs in RA in relation to traditional DMARDs, using two willingness-to-pay thresholds to assess cost-effectiveness: Can \$50,000 and Can \$100,000 per cost per quality-adjusted life year (QALY) gain. At a willingness-to-pay threshold of Can \$50,000 per QALY gain the results can be summarised as follows: a) in patients with early RA who have never received methotrexate (MTX), starting with biologic drugs is not cost-effective compared to the use of MTX; b) in patients with RA

in whom MTX mono-therapy was not effective, the use of biologics in association with MTX was cost-effective compared to continuing with MTX alone; c) in patients in whom MTX combination therapy or sequential DMARDs administration was not effective, the use of biologics was not cost-effective. On the other hand, at a willingness-to-pay threshold of Can \$100,000 per QALY gain, the results may be outlined thus: 1) the use of biologics in patients who have not previously undergone traditional DMARD treatment is only slightly cost-effective; 2) in patients in whom MTX in mono-therapy was not effective, the use of biologics was cost-effective; 3) in patients who had no response to treatment with MTX in combination with (an)other DMARD(s), introduction of the biologic drug into the treatment programme was cost-effective in 14 out of 35 comparisons. As regards other biologic drugs used in the treatment of rheumatoid arthritis, abatacept has been estimated to be cost-effective from the societal perspective in patients who do not respond to MTX and/or to the first TNF $\alpha$  inhibitor [42], [43], [44] and [45]. This is also true for rituximab when it is utilised in patients who fail to respond to the first TNF $\alpha$ , compared to another anti TNF $\alpha$  or a sequence of TNF inhibitors [43], [44], [45] and [46]. Rituximab seems to be the most cost-effective second line biologic drug [43], [44], [47] and [48]. A recent analysis has demonstrated that the use of tocilizumab is cost-effective if given after the failure of another biological DMARD or as an addition to the current standard of care in the treatment of RA with an inadequate response to DMARDs [49]. We have not found any pharmacoeconomic studies on golimumab or on certolizumab pegol. The drafting of the NICE guidelines [44] and EULAR criteria [42] and [50] for the management of RA also took into account the results of the pharmacoeconomics surveys on the cost-effectiveness of biologics.

Until the cost of biological drugs drops, the challenge is to optimise their use. This can be done through the early treatment of patients who do not respond to traditional DMARDs [39], as well as by identifying the group of patients in whom biologics can be successfully discontinued after a reasonable time without subsequent relapse of disease [11], or by identifying the subjects whose disease activity can be kept low by administering traditional DMARDs alone after the biologics. Indeed, the use of biologics after the MTX failure is cost-effective and there is evidence that these drugs can be successfully discontinued [51] and [52]. From an analysis of the BeSt study we evince that 51% of 115/508 patients achieved drug-free remission for a median duration of 23 months in the 5-year period [51]. In a study regarding a small number of cases, it was found that 42% of patients maintain low disease activity using A-Cyclosporin and MTX combination therapy: this low disease activity was obtained after 6–8 months with TNF  $\alpha$  inhibitors and MTX [52]. Predictive factors of drug-free remission in patients with RA treated with biologics have yet to be precisely defined [53], although it can be hypothesised that such factors may be sought among the predictive factors for remission found in RA patients on anti-TNF drugs such as a lower baseline HAQ score, the concurrent use of NSAIDs, a higher number of previous DMARDs, RF negativity, age at baseline and male gender [54], [55] and [56].

Besides early treatment of RA to optimise the use of biologics, tight control of the disease is important [57], [58] and [59]. Indeed, this makes it possible to more easily achieve good clinical response,

remission, prevention of anatomical lesions and swifter recovery of the functional limitations in a short time. This can lead to a decrease in the use of health resources, and most markedly, in a reduction in expenses for surgical orthopaedic treatment, together with the possibility to discontinue the administration of biological drugs sooner. It would also be possible to have a reduction in the indirect costs involved in loss of work productivity.

It is interesting to highlight that the trend towards reducing the orthopaedic treatment of RA began before the use of biological drugs and coincided with the importance that the rheumatology community attributes

to early diagnosis and early use of DMARDs [11], [60] and [61]. In the context of countering disease activity early, and in order to achieve remission as soon as possible, thus saving health resources and permitting an earlier return to work, it is necessary to assess whether among the available biological drugs there are any faster acting drugs against the inflammatory process that can lead to swifter remission of disease activity. One such candidate might be tocilizumab in view of its rapidity in suppressing the inflammatory response [62] and in producing early inhibition of bone homeostasis [63]. Rapid improvement in the signs and symptoms of rheumatoid arthritis following certolizumab pegol treatment was also shown to predict better long term outcomes in RA [64] and [65]. These elements might have a significant impact on earlier clinical improvement, prevention of joint damage and swifter recovery of function.

Considering the efficacy of synthetic DMARDs when they are used with a tight control strategy, together with recent evidence of how early response (within 1–3 months) to treatment with biological drugs is predictive of the response at 12 months, it would be interesting to know the effectiveness of tight control strategies utilising prompt cycling and swapping among various biological drugs.

As regards treatment side effects, it would also be useful to know whether there is any variability among the various biologics which might influence their cost-effectiveness. There might be a difference among TNF inhibitors: for instance, the data of the RATIO registry [66] and of the British Society for Rheumatology Biologics Register [67] reveal a similar incidence for opportunistic infections [66] and for severe infections [67], but a higher incidence of tubercular infections in patients treated with infliximab and adalimumab compared to etanercept. To answer these questions exhaustively, it would be important to conduct controlled head-head studies among the various biological drugs, also assessing possible associations with other drugs like DMARDs and steroids which could themselves increase the risks of infections. In particular, respiratory infections are among the most common causes of morbidity in the elderly and require considerable health resources, as well as also being a leading cause of death in RA. Respiratory infections have also been linked to the use of steroids and not only to TNF inhibitors and MTX [68] and [69]

In conclusion, in order to achieve the best cost-effectiveness in the treatment of RA in clinical practice, the factors that currently seem to play an important role are speed of action of the drug, tight control in patient management, and careful assessment of the safety profile of the various drugs.

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